FDA 510(k) Premarket Notification

K912165 LANDMARK DUAL-LUMEN MIDLINE CATHETER SYSTEM

CLEARED

Dec 11, 1991
209 days to decision

K912165 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k912165/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 16, 1991
Decision date	Dec 11, 1991
Days to decision	209 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Menlo Care, Inc.**
Location	Menlo Park, CA, US
Contact	JOCK M WALKER
510(k) history	31 submissions · 25 cleared · 1986-1997